Claim Amendments

Claim 1 (currently amended): A method for analyzing a human cell having an average straight line velocity of between 0 and 10 µm/min curvilinear velocity of less than 8 µm/min by suppressing movement of the human cell caused by other than activity of the human cell itself comprising the steps of:

placing the human cell having an average straight line velocity of between 0 and $10 \mu m/min$ curvilinear velocity of less than $8 \mu m/min$ in a solution containing a viscosity enhancement medium; and

measuring the motility of the human cell in the solution.

Claim 2 (original): A method as described in Claim 1 wherein the viscosity enhancement medium is methyl cellulose.

Claim 3 (previously presented): A method as described in Claim 1 wherein the viscosity enhancement medium is hyaluronic acid or chondroitin sulfate or cellulose ester or polysaccharide.

Claim 4 (original): A method as described in Claim 1 wherein multiple cells are measured in parallel.

Claim 5 (original): A method as described in Claim 1 wherein the placing step includes the step of placing the cell in the solution of between 0.1% to 0.2% by total volume of methyl cellulose for 2D analysis of motility.

Claim 6 (original): A method as described in Claim 2 wherein the placing step includes the step of placing the methyl cellulose solution having a concentration of between 0.1% and 1.2% methyl cellulose onto cells in culture medium to provide a layer of methyl cellulose-containing medium for 2D analysis of motility.

Claim 7 (original): A method as described in Claim 6 wherein the placing step includes the step of placing the cell in the solution having a viscosity of 100-5000 centipoise.

Claim 8 (original): A method as described in Claim 1 wherein the placing step includes the step of placing cells in solution having a concentration of between 0.3% to 2.5% weight per volume methyl cellulose for analysis of motility in 3D.

Claim 9 (currently amended): A method for analyzing a human cell by suppressing movement of the human cell caused by other than activity of the human cell itself comprising the steps of:

placing the cell having an average straight line velocity of between 0 and 10 μm/min curvilinear velocity of less than 8 μm/min in a solution; and

measuring the motility of the human cell in the solution when there is no attachment of the cell to any surface of the cell involved.

Claim 10 (canceled)

Claim 11 (currently amended): A method for analyzing a human cell comprising the steps of:

placing the human cell having an average straight line velocity of between 0 and $10 \mu m/min$ curvilinear velocity of less than 8 $\mu m/min$ in a solution having a viscosity of about 100-5000 centipoise; and

performing two-dimensional or three-dimensional migration analysis on the cell in the solution.

Claim 12 (currently amended): A method for analyzing a cell comprising the steps of:

placing the cell in a solution having a viscosity of about 100-5000 centipoise; and

analyzing migration of the cell in the solution which occurs without adherence of the cell to any surface.

Claims 13-15 (canceled)

Claim 16 (original): A method as described in Claim 1 wherein the placing step includes the step of placing the cell in the solution of between 1% to 5% by total volume of methyl cellulose and a concentration of between 0.08% and 0.12% of methyl cellulose.

Claim 17 (currently amended): A method for analyzing a human cell comprising the steps of:

placing the human cell having an average straight line velocity of between 0 and $\frac{10 \mu m}{min}$ curvilinear velocity of less than 8 $\mu m/min$ in a solution having a viscosity of about 100-5000 centipoise; and

measuring motility of the cell in the solution, where surface attachment by the cell to any surface is not utilized.

Claims 18-22 (canceled)

Claim 23 (currently amended): A method for analyzing a human cell by suppressing movement of the human cell caused by other than activity of the human cell itself comprising the steps of:

placing the human cell having an average straight line velocity of between 0 and 10 μm/min curvilinear velocity of less than 8 μm/min in a solution; and

placing methyl cellulose in the solution to reduce ambient motion of the human cell in the solution and eliminate convective motion.

Claim 24 (canceled)

Claim 25 (currently amended): A method for analyzing a human cell having a velocity of less than 50 µm/min by suppressing movement of the human cell relative to its location on a plate caused by forces other than activity of the human those generated by the cell itself comprising the steps of:

placing the human cell in a solution; and

using methyl cellulose in the solution for <u>suppressing a tendency for the cell to</u>
move downward on sloped surfaces of a plate holding the solution in which the cell is disposed
stopping the effects of gravity on the human cell in the solution.

Claim 26 (currently amended): A method for analyzing a human cell by suppressing movement of the human cell caused by other than activity of the human cell itself comprising the steps of:

placing the human cell having an average straight line velocity of between 0 and 10 μm/min curvilinear velocity of less than 8 μm/min in a solution; and

using methyl cellulose in the solution for reducing or eliminating the effects of micro-turbulances due to thermal convection in the solution.

Claim 27 (currently amended): A method for analyzing a human cell comprising the steps of:

placing the human cell having an average straight line velocity of between 0 and 10 μm/min curvilinear velocity of less than 8 μm/min in a solution; and

introducing methyl cellulose in the solution for stopping motion of the eells cell due to movement of the solution induced by mechanical movement of a plate on which the cells are disposed.

Claim 28 (currently amended): A method for analyzing a human cell comprising the steps of:

placing the human cell having an average straight line velocity of between 0 and 10 μm/min curvilinear velocity of less than 8 μm/min in a solution; and

introducing a viscous fluid having a viscosity of about 100-5000 centipoise in the solution for stopping or reducing the effects of gravity on the cell.

Claim 29 (currently amended): A method for analyzing a human cell comprising the steps of:

placing the human cell having an average straight line velocity of between 0 and 10 μm/min curvilinear velocity of less than 8 μm/min in a solution; and

introducing a viscous fluid having a viscosity of about 100-5000 centipoise in the solution for reducing the effects of micro-turbulences due to thermal convection.

Claim 30 (currently amended): A method for analyzing a human cell comprising the steps of:

placing the cell in a solution; and

introducing a viscous fluid having a viscosity of about 100-5000 centipoise in the solution for stopping motion of the eells cell due to effects on the cell of currents in the solution that are induced by motion of a mechanical movement of the plate on which the cell is disposed.

Claim 31 (currently amended): A method for analyzing a human cell by suppressing movement of the human cell caused by other than activity of the human cell itself comprising the steps of:

placing the human cell having an average straight line velocity of between 0 and 10 μm/min curvilinear velocity of less than 8 μm/min in a solution; and

using methyl cellulose or any viscous fluid to separate biological motility from ambient motility.

Claims 32-37 (canceled)

Claim 38 (previously presented): A method for analyzing either a T-cell, dendritic cell, B-cell or lymphocyte having an average straight line velocity of between 0 and 10 µm/min by suppressing movement of either a T-cell, dendritic cell, B-cell or lymphocyte caused by other than activity of either a T-cell, dendritic cell, B-cell or lymphocyte itself comprising the steps of:

placing either a T-cell, dendritic cell, B-cell or lymphocyte having an average straight line velocity of between 0 and 10 μ m/min in a solution containing a viscosity enhancement medium; and

measuring the motility of either a T-cell, dendritic cell, B-cell or lymphocyte in the solution.